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Short communication

Study of disease-relevant polymorphisms in the *TLR4* and *TLR9* genes: A novel method applied to the analysis of the Portuguese population

A. Carvalho^a, A. Marques^b, P. Maciel^a, F. Rodrigues^{a,*}^a*Life and Health Sciences Research Institute (ICVS), School of Health Sciences, University of Minho, Braga, Portugal*^b*Department of Immunohemotherapy, Hospital de São Marcos, Braga, Portugal*

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Abstract

Toll-like receptors (TLRs) are cellular receptors that mediate recognition of microbial challenges and the subsequent inflammatory response. Genetic variations within these inflammation-associated genes may alter host-pathogen defence mechanisms affecting susceptibility towards infectious diseases. Taking into account the significance of these genes, we developed a simple and rapid method based in the bi-directional PCR amplification of specific alleles (Bi-PASA) for genotyping known sequence variants in *TLR4* (Asp299Gly and Thr399Ile) and *TLR9* (T-1237C) genes. This method allows genotype determination in a single reaction and is amenable to large-scale analysis. We used Bi-PASA to characterize the distribution of these polymorphisms in the Portuguese population. A total of 388 randomly selected blood donors of Portuguese origin (203 females and 185 males) were genotyped and allele frequencies were determined. Among the tested individuals, 11.1% and 10.8% were heterozygous for Asp299Gly and Thr399Ile, respectively. In what concerns the T-1237C variation in *TLR9*, the variant allele was present in 19.4% of the individuals tested. Besides confirming the usefulness of the Bi-PASA in polymorphism analysis, the data presented provide valuable information on *TLR* polymorphisms in the Portuguese population that can be used to stratify risk patients with increased susceptibility to infection.

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1. Introduction

The innate immune system is able to recognize conserved motifs in pathogens in which pattern recognition receptors, including toll-like receptors (TLRs), play an important role [1]. TLRs are a family of genetically conserved proteins identified as key components of the innate immune system, mediating recognition of microbial challenges and the subsequent inflammatory response [2]. The first of the currently known TLRs was described in 1997 as a human homolog of the *Drosophila* Toll protein, later designated *TLR4* [3]. The complete TLR family allows the host to detect infection by most, if not all, types of microbial pathogens. Well characterized receptor–ligand pairs include *TLR4* and lipopolysaccharide (LPS) [4,5], *TLR5* and bacterial flagellin [6], *TLR3* and viral double-stranded

DNA [7], *TLR9* and bacterial hypomethylated DNA [8] and *TLR2*, in association with *TLR1* or *TLR6*, and a variety of cell wall components from gram-positive bacteria [9].

Cellular activation via TLRs triggers not only innate immune responses but also initiates adaptive immunity [10]. Due to the significance of TLRs in the immune system, genetic variations within these genes could have a major impact upon host immune response to pathogens and thus, an increased susceptibility to infection. Regarding TLRs, several polymorphisms are already largely studied and characterized concerning the phenotypical outcome, such as polymorphisms in *TLR4* [A+896G (SNP ID: rs4986790) and C+1196T (SNP ID: rs4986791)]. These polymorphisms are located in the coding sequence resulting in amino acid exchanges: an aspartic acid for a glycine at position 299 (Asp299Gly) and a threonine for an isoleucine at position 399 (Thr399Ile), respectively, affecting the extracellular domain of this receptor. Others, such

*Corresponding author. Tel.: + 351 253 604814; fax: + 351 253 604809.
E-mail address: fr Rodrigues@ecsau.de.uminho.pt (F. Rodrigues).

Table 1

Summary of association studies between *TLR4/TLR9* polymorphisms and infectious and inflammation-related diseases

Reference	Association study	Patients (<i>n</i>)	Controls (<i>n</i>)	TLR SNP		Case vs. control (%)	<i>p</i> value
Arbour et al. [11]	Hyporesponsiveness to inhaled lypopolysaccharide	31	57	TLR4	Asp299Gly	22.6 vs 5.8	0.029
Lorenz et al. [13]	Septic shock	91	73		Thr399Ile		
Lorenz et al. [32]	Premature birth associated to infection	440	351		Asp299Gly	5.5 vs 0	0.05
					Asp299Gly	23.8 vs 15.9	0.028
Agnese et al. [12]	Gram-negative infections	77	39		Thr399Ile		
					Asp299Gly	18.0 vs 12.8	0.004
Tal et al. [31]	Respiratory syncytal virus infection	99	82		Thr399Ile		
					Asp299Gly	20.2 vs 4.9	0.003
Edfeldt et al. [33]	Myocardial infarction associated to infection	1213	1561		Thr399Ile		
					Asp299Gly	10.7 vs 7.9	0.004
Hawn et al. [34]	Legionnaire's disease	108	510		Thr399Ile		
					Asp299Gly	2.5 vs 6.5	0.025
Rezazadeh et al. [35]	Brucellosis	198	111		Thr399Ile		
Mockenhaupt et al. [36]	Severe malaria	70	51		Asp299Gly	33.6 vs 20.7	<0.0001
					Thr399Ile	24.1 vs 17.6	<0.05
Lazarus et al. [17]	Asthma	67	152	TLR9	T-1237C	23.4 vs 14.5	0.042
Torok et al. [18]	Crohn's disease	174	265		T-1237C	19.3 vs 11.9	0.0036

as those concerning *TLR9*, have just recently started to be characterized and their functional importance elucidated. Among these, the most studied is T-1237C (SNP ID: rs5743836), a polymorphism located within the putative promoter region that may influence transcriptional regulation of the *TLR9* gene.

Arbour and colleagues were the first to describe that the two mutations affecting the extracellular domain of TLR4 protein at the 299 and 399 residues were associated with blunted physiological responses to inhaled lipopolysaccharide [11]. A large number of association studies followed and the most relevant are summarized in Table 1. Since their identification, these two mutations have been studied for their association with various infectious and inflammatory diseases. Specifically, a positive association of the Asp299Gly mutation with increased susceptibility to Gram-negative bacteremia and septic shock was found [12,13]. Interestingly however, no association between the Asp299Gly and/or Thr399Ile mutations was found for other diseases of infectious etiology, including candidiasis [14], tuberculosis [15] or meningococcal disease [16]. The variation T-1237C in the promoter region of *TLR9*, on the other hand, has been shown to be associated with an increased risk for asthma [17] and preliminary data regarding a possible positive association with Crohn's disease was also reported [18]. Nevertheless, some studies also report no association between this polymorphism and susceptibility to systemic lupus erythematosus [19] and atopy [20].

Thus, assessment of polymorphisms in TLRs may have a potential clinically usefulness for risk stratification of

patients possibly more vulnerable to infections. Some of the previously developed methods are, to a certain extent, attractive laboratory-tailored approaches for genotyping polymorphisms in *TLR* genes, such as the real-time PCR-based method developed by Hamann et al. [21] or the single tube PCR reaction based on exonuclease degradation of allele-specific probes described by van Rijn et al. [22]. However, most of the genotyping techniques currently available are still either time-consuming and laborious or require expensive equipment [23–25]. In this sense, we adapted a simple and rapid method, allowing genotype determination in a single reaction based in the bi-directional PCR amplification of specific alleles (Bi-PASA) [26] in both the *TLR4* (Asp299Gly, Thr399Ile) and *TLR9* (T-1237C) genes, a methodology amenable to large scale analysis, using technology available in most scientific laboratories.

2. Materials and methods

Bi-PASA uses a combination of four primers, two outer primers (named P and Q) and two inner allele-specific primers (termed M and W). The inner primers are characterized by containing a 10-nucleotide G + C-rich 5' tail, which prevents “megapriming” and enhances the efficiency of amplification. Depending on the genotype, Bi-PASA produces two or three overlapping fragments. PQ is always produced and serves as a positive control. PW and MQ are present in a heterozygote individual, but PW is only produced in wild-type homozygote and MQ only in

homozygous mutant samples (Fig. 1). Primers for Bi-PASA were designed following the guidelines proposed by Liu et al. [26], taking into account both the melting temperature of both the primers and the largest PCR segment (PQ). The primers used in this study are shown in Table 2. The Bi-PASA methodology was optimized using DNA samples from individuals with known genotype for the polymorphisms being studied (Fig. 1(B)–(D)). These samples were previously genotyped by Berghöfer et al. and Hamann et al., using real-time PCR-based methodologies and confirmed by direct sequencing [20,21].

DNA was isolated from whole blood samples using the salting-out procedure described by Miller et al. [27] and PCR amplification was performed in a 20 µl volume that included autoclaved ultra-filtered water, PCR buffer (1.6 ×), dNTP mixture (200 µM each), primers (0.05–0.4 µM each; for details see Table 2), *Taq* DNA polymerase (1 U/20 µl) and approximately 50 ng genomic DNA templates. PCR cycling conditions included 35 cycles of 15 s at 94 °C, 30 s at 57 °C, and 45 s at 65 °C, after a

10-min initial period of DNA denaturation and enzyme activation at 94 °C. The amplified fragments had sizes readily distinguishable by electrophoresis through a 2% agarose gel. The results obtained for the genotype of the Portuguese population by Bi-PASA were further validated by sequencing analysis of 10 randomly selected samples (data not shown).

3. Results and discussion

After written informed consent was obtained from each volunteer, a total of 388 randomly selected blood donors of Portuguese origin (203 females and 185 males) were genotyped using Bi-PASA. Among the tested individuals, 43/388 (11.1%) and 42/388 (10.8%) were heterozygous for Asp299Gly and Thr399Ile polymorphisms, respectively. None of the individuals showed a homozygous *TLR4* polymorphism. In what concerns the T-1237C variation in *TLR9*, 67/388 (17.3%) were heterozygous and 8/388 (2.1%) were homozygous for this polymorphism. Based

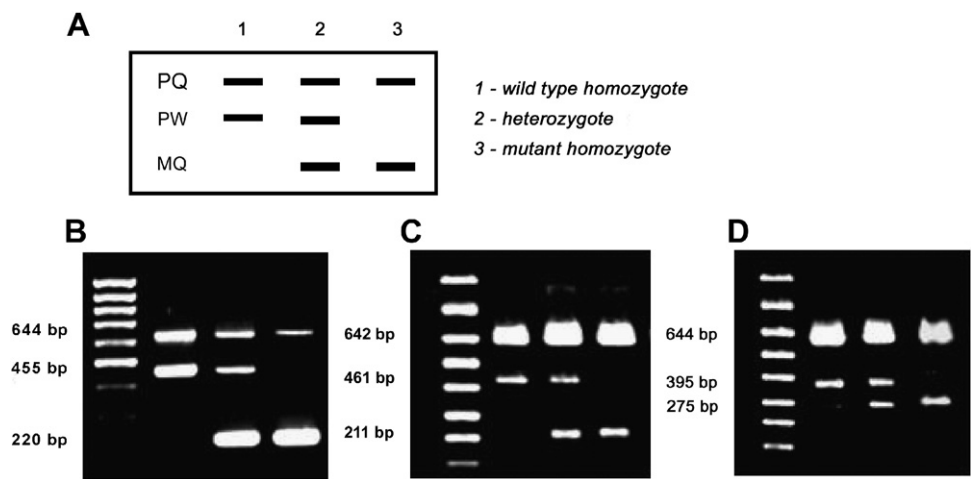


Fig. 1. Predicted band pattern obtained with Bi-PASA genotyping (A). Agarose gel electrophoresis showing results of Bi-PASA genotyping for Asp299Gly (B), Thr399Ile (C) and T-1237C (D).

Table 2
Bi-PASA primers used in this study

Gene	Polymorphism	Primer	Primer final concentration (µM)
TLR4	Asp299Gly	P: 5'-AGAACTTAATGTGGCTCACAAT-3'	0.1
		Q: 5'-GAAAAAGCATTCCCACCTTTG-3'	0.1
		W: 5'-gggccgggggTACTACCTCGATGG-3'	0.4
		M: 5'-ggcgccggggTTAAATAAGTCAATAATAT-3'	0.4
	Thr399Ile	P: 5'-CTGGCTGGTTTAGAAGTCCA-3'	0.1
		Q: 5'-ATTGAAAGCAACTCTGGTGTG-3'	0.1
		W: 5'-gggccgggggAAATACTTTAGGCTGT-3'	0.1
		M: 5'-ggcgccggggTGATTTTGGGACAAT-3'	0.1
TLR9	T-1237C	P: 5'-TCATTAGCCTTCACTCAGA-3'	0.4
		Q: 5'-CACATTCAGCCCCTAGAGGG-3'	0.4
		W: 5'-ggcgccggggGTGCTGTTCCCTCTGCCTGA-3'	0.05
		M: 5'-gggccgggggATGAGACTTGGGGGAGTTTC-3'	0.05

Table 3

Allele frequencies of each *TLR* polymorphism in distinct ethnic populations [data from Entrez SNP database (<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?CMD=search&DB=snp>) in November, 2006] and in our study in the Portuguese population

TLR SNP		Allele frequencies				
		This study	European	Asian	African American	Sub-Saharan African
Asp299Gly	A	0.889	0.967	1.000	0.848	0.967
	G	0.111	0.033	0	0.152	0.033
Thr399Ile	C	0.892	0.967	1.000	n/a	1.000
	T	0.108	0.033	0		0
T-1237C	C	0.806	0.841	0.974	0.750	n/a
	T	0.194	0.159	0.026	0.250	

n/a—not available.

on the control samples of a large number of published studies [28], the two *TLR4* polymorphisms, Asp299Gly and Thr399Ile, are known to be present at an overall allele frequency of 6.4 ± 2.8 (range, 0–19.6; $n = 38$ studies) and 7.2 ± 3.8 (range, 0–13, $n = 11$), respectively. However, reports of differences in the frequencies of these two polymorphisms among different ethnic groups have been published [15,29]. In fact, considering the data currently available on the Entrez SNP database (Table 3), the frequency of *TLR4* Asp299Gly polymorphism is much higher in African Americans than in Europeans and sub-Saharan Africans, in contrast to Asians, where this particular polymorphism does not seem to occur. The overall frequencies of Thr399Ile are similar to that of Asp299Gly, although no data is available for African Americans. African Americans also display a higher frequency of the *TLR9* polymorphism than Europeans and Asians. Comparing with our results, we can observe that *TLR4* Asp299Gly and Thr399Ile allele frequencies in the Portuguese population are higher than those described for the European population characterized in the Entrez database, although similar to those of controls published in association studies using European populations. Concerning *TLR9*, the frequency of T-1237C in the Portuguese population is similar to the one presented in the database for the European population. Distributions of both *TLR4* and *TLR9* genotypes did not deviate from those predicted by the Hardy-Weinberg equilibrium (Asp299Gly, $p = 0.388$; Thr399Ile, $p = 0.410$; T-1237C, $p = 0.58$). In addition and as expected, both *TLR4* polymorphisms are in total linkage disequilibrium ($D' = 1.000$). In fact, cosegregation among these two mutations has also been largely stated [13,30,31].

In conclusion, the data herein presented provides valuable information on *TLR* polymorphisms in the Portuguese population that can be used in future studies to stratify patients regarding susceptibility to infection. Our results clearly show that Bi-PASA is a valuable methodology for genotyping studies. In addition, this method can easily be applied in studies of large populations, and further applied for the study of other polymorphisms, as it is a cost-effective technique, where in a

single reaction the genotype of the individuals can be determined.

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